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Publication number:

**0 083 494  
A1**

## EUROPEAN PATENT APPLICATION

Application number: 82306834.1

Int. Cl.: **A 61 F 9/00**

Date of filing: 21.12.82

Priority: 04.01.82 US 336919

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Date of publication of application: 13.07.83  
Bulletin 83/28

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Designated Contracting States: DE FR GB IT

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Apparatus for corneal curvature adjustment.

An apparatus for adjusting the curvature of the cornea of the eye comprises a plastic split end adjusting ring which is insertable into the stroma of the cornea and a metal split end dissecting ring, one end of which is insertable into the cornea with the trailing end of the dissecting ring remaining outside the cornea. The one end of the dissecting ring has a transverse hole therein near its tip end. The dissecting ring is held by a magnetic holding and rotating device which has a concave end surface and circular groove therein for receiving the metal dissecting ring and for holding it in a circular shape. As the holder is rotated, the inserted end of the metal ring is magnetically forced to follow rotation of the holder, thus the dissecting ring is displaceable, in use, in a circular path within the stroma. A sled shaped end portion of the dissecting ring permits, in use, the moving inserted end of the dissecting ring to be biased upwardly toward the anterior of the stroma as the dissecting ring is rotatably inserted therein. The adjusting member also has a transverse hole near its one sled shaped end which is placed next to the first incision. A connecting link member is inserted through the holes in the ends of the respective dissecting and adjusting rings to releasably join the two rings, the rotational direction of the holding tool can then be reversed to permit, in use, the dissecting ring to be removed and at the same time this pulls in the plastic adjusting ring.

ACTORUM AG

APPARATUS SUITABLE FOR CORNEAL CURVATURE ADJUSTMENTBACKGROUND OF THE INVENTION

This invention relates to apparatus suitable for adjusting the shape of components of the eye and more particularly to making fixed changes in the corneal curvature.

5           Deviations from the normal shape of the corneal surface produce errors of refraction in the visual process. The eye in a state of rest, without accommodation, focuses the image of distant objects exactly on the retina. Such an eye enjoys distinct vision for distant objects without effort. Any varia-  
10       tion from this standard constitutes ametropia, a condition in which the eye at rest is unable to focus the image of a distant object on the retina. Hyperopia is an error of refraction in which, with the eye at rest, parallel rays from distant objects are brought to focus behind the retina. Divergent rays from  
15       near objects are focused still further back. In one aspect of hypertopia, the corneal surface is flattened which decreases the angle of refraction of rays as they pass through the refractive surfaces of the cornea, causing a convergence or focus of the rays at a point behind the retina. The  
20       retina is comprised partially of nerve fibers which are an expansion of the optic nerve. Waves of light falling on the retina are converted into nerve impulses and carried by the optic nerve to the brain to produce the sensation of light. To focus parallel rays on the retina, the hyperopic eye must  
25       either accommodate, i.e., increase the convexity of its lens, or a convex lens of sufficient strength to focus rays on the retina must be placed before the eye.

30       Myopia is that refractive condition in which, with accommodation completely relaxed, parallel rays are brought to focus in front of the retina. One condition which commonly causes myopia is when the corneal curvature is steepened, thus the refraction of rays is greater as they pass through the refractive surfaces of the cornea, and the over refracted rays

converge or focus in front of the retina in the vitreous of the eye. When the rays reach the retina they become divergent, forming a circle of diffusion and consequently a blurred image. A concave lens is used to correct the focus of the eye for myopia.

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The normal treatment of these classic forms of refractive error of the eye is with the use of eyeglasses or contact lenses, both of which have well-known disadvantages to the user. Recent research has been directed to operative techniques to change the refractive condition of the eye. Such techniques are generally referred to "keratorefractive techniques". Two such techniques are more particularly called keratophakia and keratomileusis. Keratomileusis involves the regrinding of a corneal lamella into a meniscus or hyperopic lens to correct myopia or hyperopia. A corneal optical lathe has been especially developed for this procedure and is also used in the keratophakia procedure, when a homograft ground into a convex lens is placed interlamellarly to correct aphakic hypermetropia. The homograft tissue (corneal lamella) is frozen with carbon dioxide. The homograft is cut as a contact lens would be, i.e. to the optical power required to effect the desired optical correction of the cornea. In keratomileusis, the anterior corneal lamella is shaped by the lathe and in keratophakia, it is the corneal stroma of a donor eye that is shaped by the lathe. These techniques have a broad application in the correction of high hyperopic and myopic errors. These procedures require radial cutting of the cornea about the periphery of the graft which weakens the cornea so that pressure from fluids below the incisions pushes up under the cuts and flattens the curvature of the cornea. This flattening of the cornea results in refractive errors to the eye not compensated for by the graft. Suturing in these operations also causes radial asymmetry of the cornea consequently promotes astigmatic error in this regard. Sutures also cause scarring of the corneal tissue, which scar tissue loses its transparency. Surgical correction of

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BAD ORIGINAL

astigmatism is accomplished by asymmetrically altering the corneal curvatures. The effect of a peripheral distorting force may be easily visualized by imagining an inflated balloon with a spherical surface being compressed between the palms of the hands. Because the volume of air in the balloon is constant, the surface area remains constant. The previously spherical anterior surface is distorted meridianally as a result of compressing the diameter between the hands so that the curvature changes without changing the circumference of the surface. The meridian passing over the balloon between the extended fingers steepens, while the uncompressed meridian at right angles thereto flattens as its diameter lengthens in proportion to the shortening of the compressed diameter. This demonstrates the effect that may result from slight variations in the symmetrical patterns or intentional asymmetrical patterns attempted to be accomplished during surgical procedures and attendant suturing. It is thus seen that present procedures in keratorefractive techniques are best limited to situations where other more standard corrective practices are found ineffective. It is readily seen that the limiting factors in such surgical techniques is the gross complexity involved not only with multiple incisions in corneal tissue for affecting the procedures but also complex suturing patterns, resulting in gross restructuring of the eye. The eye is thus faced with a difficult job of adjusting to this trauma.

It is therefore an object of the present invention to provide a new and improved keratorefractive technique involving apparatus for changing the shape of the optical zone of the cornea to correct refractive error whereby a minimum disturbance is imposed on the eye system and the simplicity of the technique virtually eliminates the chance of error or further complications resulting from gross disturbances of the eye system.

SUMMARY OF THE INVENTION

5 With this and other objects in view of the present invention contemplates apparatus involving including a split end adjusting ring, one end of which is insertable in the cornea of the eye, the ring then being moved in a circular path until its ends meet. Thereafter the ends are adjustable  
relative to one another until the shape of the eye has assumed a desired curvature whereupon the ends are fixedly attached to one another maintain the desired curvature of the cornea.

10 Preferably, the cross sectional shape of the adjusting ring is ovaloid or ellipsoid such that, when inserted in the cornea, its major cross sectional axis is alignable with a corneal arc extending through the anterior pole of the  
15 cornea.

Another aspect of the invention provides a dissecting ring which has split end portions, one end thereof being insertable into the cornea prior to insertion of the adjusting  
20 ring. Such one end is movable in a circular path about the interior of the cornea until it reaches the insertion point, whereupon the one end of the adjusting ring is releasably attachable to the one end of the dissecting ring. The dissecting ring can then be moved in a reversed circular path,  
25 thereby pulling the now attached adjusting ring behind it until the one end of the dissecting ring has returned to the insertion point. At this time, the one end of the adjusting ring has also circularly moved about the interior of the cornea until its one leading end has reached the insertion  
30 point to implant the adjusting ring in the cornea and withdraw the dissecting ring. The corneal curvature adjusting procedure and ring end fixing procedure are then performed.

Still another aspect of the invention is the shape of the one leading end of the respective dissecting ring and adjusting ring which is asymmetrically rounded into a sled shape. In use, this maintains a transverse bias on the one ends during movement thereof within the corneal tissue.

Yet another aspect of the invention pertains to a connecting system for releasably attaching the leading ends of the respective dissecting and adjusting rings, including holes near the tip ends of the one end of the rings with longitudinal grooves extending from the holes to the tip ends, and an inverted "U-shaped" clip which is inserted in the holes of both rings to hold the ring ends together while they are being moved about the interior of the cornea.

Another aspect of the invention involves the use of a ring holder and moving device which comprises an elongated cylindrical member having a concave surface formed concentrically within its lower end and sized so as, in use, to fit over the curvature of the cornea, with a circular groove in the face of the concave surface, which groove is sized to hold the dissecting ring in its circular configuration, and a means for exerting a magnetic force on the dissecting ring to cause the ring to follow the rotational path of the holder when it is rotated over the eye after the one end of the dissecting ring is inserted into the corneal tissue. This magnetic force also pulls, in use, the dissecting ring toward the anterior of the corneal stroma into which it is being inserted. In addition, a pin extends outwardly from the groove and serves to engage a blunt trailing end of the dissecting ring as it is being inserted to further facilitate its movement in and about the cornea as the holder is rotated. Also, a cylindrical sleeve may be fitted over a portion of the elongated cylindrical member to facilitate holding the holding device

steady as the cylindrical member is rotated in the adjusting ring insertion procedure.

5 Yet still another aspect of the invention resides in providing visual indication of the present shape of the cornea together with an indication of a desired shape and comparing the indications while the adjusting ring ends are being adjusted to aid in fixing the ends of the adjusting ring at a place to provide the desired corneal topography.

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#### BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic illustration of a horizontal section of the eye;

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Fig. 2 is a schematic illustration of an eye system showing adjustment of the cornea to steepen the corneal slope;

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Fig. 3 is a schematic illustration of an eye system showing adjustment of the cornea to flatten the corneal slope;

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Fig. 4 is a more detailed schematic illustration of a horizontal section of the frontal portion of the eye showing an adjusting ring positioned in the stroma of the cornea;

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Fig. 5 is a plan view of a dissecting ring showing its end portions;

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Fig. 6 is a plan view of an adjustment ring showing its end portions;

Fig. 7 is an elevational view showing the inserting ends of the respective dissecting ring and adjustment rings positioned for receiving a releasable connecting clip;

Fig. 8 is an elevational view of the adjusting ring taken along lines 8-8 of Fig. 6;

Fig. 9 is a partial perspective view of the cornea of an eye with incisions for receiving the dissecting and adjusting rings and the rings positioned for releasable attachment to one another just prior to implanting the adjusting ring;

Fig. 10 is a side elevational cross sectional view of a dissecting ring holding and rotating tool; and

Fig. 11 is a schematic representation of a cornea-scope type image superimposed with a target image for comparing the present shape of the cornea with a desired shape to permit accurate fixing of the adjustment ring to fix the shape of the cornea.

#### DESCRIPTION OF THE PREFERRED EMBODIMENT

Referring first to Fig. 1 of the drawings, a horizontal section of the eye shows the globe of the eye resembling a sphere with an anterior bulged spherical portion representing the cornea. Thus the eye is actually comprised of two somewhat modified spheres placed one in front of the other. The anterior of these two segments is the smaller more curved cornea.

The globe of the eye consists of three concentric coverings enclosing the various transparent media through which the light must pass before reaching the sensitive retina. The outermost covering is a fibrous protective portion of the posterior five-sixths of which is white and opaque and called the sclera 13, and sometimes referred to as the white of the eye where visible to the front. The anterior one-sixth of this outer layer is the transparent cornea 12.



A middle covering is mainly vascular and nutritive in function and is comprised of the choroid 14, ciliary body 15 and iris 17. The choroid generally functions to maintain the retina. The ciliary muscle is involved in  
5 suspending the lens and accommodation of the lens. The iris is the most anterior portion of the middle covering of the eye and is arranged in a frontal plane. It is a thin circular disc corresponding to the diaphragm of a camera, and is perforated near its center by a circular aperture called  
10 the pupil 19. The size of the pupil varies to regulate the amount of light which reaches the retina. It contracts also to accommodation, which serves to sharpen the focus by diminishing spherical aberration. The iris divides the space between the cornea 12 and the lens 21 into an anterior chamber  
15 22 and posterior chamber 23. The innermost portion of covering is the retina 18, consisting of nerve elements which form the true receptive portion for visual impressions.

The retina is a part of the brain arising as an out-  
20 growth from the fore-brain, with the optic nerve 24 serving as a fibre tract connecting the retina part of the brain with the fore-brain. A layer of rods and cones, lying just beneath a pigmented epithelium on the anterior wall of the retina, serve as visual cells or photoreceptors which  
25 transform physical energy (light) into nerve impulses.

The vitreous 26 is a transparent gelatinous mass which fills the posterior four-fifths of the globe. At its sides it supports the ciliary body 16 and the retina 18.  
30 A frontal saucer-shaped depression houses the lens 21.

The lens 21 of the eye is a transparent bi-convex body of crystalline appearance placed between the iris 17 and vitreous 26. Its axial diameter varies markedly  
35 with accommodation. A ciliary zonule 27, consisting of transparent fibres passing between the ciliary body 16 and lens 21 serves to hold the lens in position and enable

the ciliary muscle to act on it.

Referring again to the cornea 12, this outermost fibrous transparent coating resembles a watch glass. Its curvature is somewhat greater than the rest of the globe and is ideally spherical in nature. However, often it is more curved in one meridian than another giving rise to astigmatism. A central third of the cornea is called the optical zone with a slight flattening taking place outwardly thereof as the cornea thickens towards its periphery. Most of the refraction of the eye takes place on the surface of the cornea.

Referring to Fig. 4, a more detailed drawing of the anterior portion of the globe shows the various layers of the cornea comprising an epithelium 31. Epithelial cells on the surface thereof function to maintain transparency of the cornea. These epithelial cells are rich in glycogen, enzymes and acetylcholine and their activity regulates the corneal corpuscles and controls the transport of water and electrolytes through the lamellae of the stroma 32 of the cornea.

An anterior limiting lamina 33, referred to as Bowman's membrane, is positioned between the epithelium 31 and the substantia propria or stroma 32 of the cornea. The stroma is comprised of lamella having bands of fibrils parallel to each other and crossing the whole of the cornea. While most of the fibrous bands are parallel to the surface, some are oblique, especially anteriorly. The fibrous bands within alternate lamella are at a near right angle to bands in the adjacent lamella. A posterior limiting lamina 34 is referred to as Descemet's membrane. It is a strong membrane sharply defined from the stroma and resistant to pathological processes of the cornea.

The endothelium 36 is the most posterior layer of the cornea and consists of a single layer of cells. The limbus 37 is the transition zone between the conjunctiva 38 and sclera 13 on the one hand and the cornea 12 on the other.

Referring next to Fig. 2 of the drawings, the globe of an eye is shown having a cornea 12 with a normal curvature represented by the solid line 39. If parallel rays of light 41 pass through the corneal surface 39 of Fig. 2 they are refracted by the corneal surfaces to converge eventually near the retina 18 of the eye. The diagram of Fig. 2 discounts, for the purposes of this discussion, the refractive effect of the lens or other portions of the eye. The eye depicted in Fig. 2 is hyperopic and thus the rays of light 41 are refracted to converge at point 42 behind the retina. If a peripheral band of pressure is applied inwardly at the chord 43 of the cornea, the walls of the cornea are caused to steepen. This is because the volume of fluids within the anterior chamber 22 remains constant, thus the anterior portion of the cornea, including the optical zone (inner third of the cornea) steepens in slope to form a curvature (shown in exaggeration) following the dotted line 44. The rays of light 41 are then refracted from the steeper surface 44 at a greater angle to direct the refracted rays into focus at a shorter distance, such as directly on the retina 18.

Fig. 3 shows a similar eye system to that of Fig. 2 except that the so called normal corneal curvature of Fig. 3 causes the light rays 41 to refract into focus at a point 46 in the vitreous which is short of the retinal surface 18. This is typical of a myopic eye. If chord 43 of the cornea is expanded uniformly outwardly as shown by the arrows, the walls of the cornea are flattened. Light rays 41 refracted by the now flattened corneal surface will be refracted at a smaller angle and thus converge

at a more distant point such as directly on the retina 18.

The apparatus of the present invention is concerned with a system suitable for adjusting an annular chord of the cornea as suggested by the processes shown in Figs. 2 and 3 to thereby correct refractive errors of the eye. Again referring to Fig. 4, a ring 47, having an ovaloid cross sectional shape is shown implanted in the stroma layer of the cornea. By adjusting the diameter of such a ring in the cornea and fixing that diameter at a discrete value, the rays refracted by the cornea and other eye components can be brought to focus directly on the retina 18. Such a ring placed approximately at the 8 mm chord of the cornea provides a means for making such a corrective adjustment. Apparatus and methods for making this adjustment are hereinafter described.

Referring now to Figs. 6 and 8 of the drawings, the adjusting ring 47 is comprised of a generally circular member having split end portions 48 and 49. The ring is comprised of a material which has sufficient stiffness to maintain its generally circular shape and sufficient resiliency to permit its ends 48 and 49 to be adjusted relative to one another to thereby enlarge or decrease the normal diameter of the ring at rest. The material should have properties that render it physiologically compatible with the tissue of the cornea. Two such materials are plastic type materials sold under the trade names PLEXIGLASS and SAUFLON. The cross sectional shape of the rings is that of an oval generally dimensioned to be about 1 mm across its major axis and .2 mm across its minor axis. The one insertion end or leading end 48 of the adjusting ring is tapered asymmetrically to a rounded tip end (See Fig. 7).

As shown in Fig. 8, the major axis of the oval ring 47 is formed at an angle sloping inwardly to the center of the ring. The angular disposition of the major axis of

the ovaloid ring corresponds to the intended implantation position of the ring 47 in the cornea. The ring is implanted in the stroma 32 of the cornea as shown in Fig. 4. One function of the stroma is to transfer fluids through the eye. In order to minimize the effect of the implanted ring 47 on the transfer of fluids, the ring is positioned so that its major axis is parallel to the lamellae of the stroma. Thus the ring is implanted at a slope corresponding to the slope of a corneal arc extending through the anterior pole 20 (Fig. 1) of the cornea. This slope of the adjusting ring also corresponds to the direction of lamellae within the corneal stroma. By orienting the ring thusly, the ring may be inserted between the lamellae to produce a minimum of trauma to the eye. During the development of this procedure, a circular adjusting ring was used. It was found however that the circular ring, when expanded or contracted to adjust its size (see Figs. 2 and 3), creates a sufficient pressure between the corneal tissue and the ring, to cause the ring to cut through the tissue. By utilizing an ovaloid shape and orienting the major axis of the ovaloid as shown in Fig. 4, an enlarged surface is presented in the direction of pressure of the ring to prevent cutting of the tissue. Thus the major axis of the ring acts against the lamellae of the stroma in the direction of pressure and the minor axis is aligned with the lamellae to provide a minimum of interruption to fluid flow.

Fig. 5 of the drawings shows a dissecting ring 51 which is circular in cross section and has overlapping split end portions 52 and 53. The ends of the ring overlap approximately one half a diameter of the ring. Ring 51 may be constructed of a metallic material such as stainless steel and in any event a magnetic material for purposes to be hereinafter described. Ring 51 also is provided with an asymmetrically rounded end portion 52 having a lower tapered surface to resemble the shape of

a sled runner. The trailing end 53 of the dissecting ring 51 has a blunt end surface 54.

5 Referring now to Fig. 7 both the dissecting ring 41 and the adjusting rings have transverse substantially vertical holes 56 and 57 respectively through their ends, near the tip ends thereof. Longitudinal grooves 58 are formed in the top surface of each of the rings 47, 51 and extend from the holes 56, 57 to the respective tip ends of the rings. An inverted U-shaped clip member 61 is shown 10 positioned in Fig. 7 for reception of its downwardly projecting leg portions 62 into the holes 56 and 57 of the dissecting and adjusting rings respectively and its body portion 63 into the longitudinal grooves 58 in the 15 top end of the respective rings.

Referring to Fig. 10 of the drawings, a dissecting ring holding and rotating apparatus 71 is shown including a solid cylindrical rotating member 72 having a knurled 20 upper end surface 73. The cylindrical rotating member 72 is slip fitted with a cylindrical holding sleeve 74 to permit relative rotation between the member 72 and sleeve 74. The lower end of the rotating member 72 has a concentrically arranged concave surface 76 having its 25 concave shape as well as the proportions of the surface 76 arranged to be matingly received over the anterior corneal surface of the eye. An annular groove 77 is formed in the concave surface 76 near the peripheral edge of the rotating member 72. This groove 77 is sized 30 to receive and hold the dissecting ring at its nominal diameter for insertion into and around the cornea at the 8 mm chord of the eye as shown in Fig. 4. This chord is located within a plane across the cornea which measures approximately 8 mm in diameter. Thus the groove 77 itself 35 has an internal diameter of approximately 8 mm. The rotatable holding member 72 is constructed of a magnetized material or has a means for being magnetized. The magnetic

nature of the member 72 holds the ring 51 in place on the tool during the keratorefractive procedure to be described. A small pin 78 is vertically arranged to project downwardly from a point in the groove. This pin contacts the trailing  
5 blunt end 54 of the dissecting ring.

The method of implanting an adjusting ring 47 within the cornea of the eye will now be described: First, a determination is made, by taking optical measurements of the  
10 eye, as to what shape the cornea should have in order for that eye to operate in an optically correct manner. Fig. 11 shows a target image comprised of indicia 81. This target grid 81 may be made by reflecting light from placido rings from a standard spherical surface the same size as the eye  
15 in question and at a fixed distance. An image is made of this "correct" topographic map of the eye, referred to herein as a target or target image. The target indicia is in the form of vertical lines or grids 81 as shown in Fig. 11 which of course may be drawn in other radial  
20 meridians as well as the horizontal meridian as shown. The spacings between the grids 81 represents a topographic survey of an eye having a corrected curvature, as pertains to the specific eye in question.

25 A dissecting ring 51 is next placed in the groove 77 on the concave surface 76 of the rotatable holding member 72. The blunt end 54 of ring 51 is contacted against the pin 78 extending outwardly from the groove. The magnetic attraction between the magnetic member 72  
30 and steel ring 51 holds the ring in place on the end of the member 72. Next, working under a surgical microscope, a small (approximately 1 mm long and .2 mm deep) incision 82 (Fig. 9) is made in the cornea through the epithilium and Bowman's membrane. This incision is approximately  
35 the same size as the adjusting ring to be implanted. The leading end 52 of the dissecting ring 51 is then moved

through the incision and into the stroma of the cornea.

The holding member 72 is then rotated to progressively thread the ring around the cornea between the adjacent lamellae within the anterior portion of the stroma. The lamellae of fibrils near the anterior of the stroma are more loosely formed, making this a desirable location in the stroma for insertion of the dissecting ring and for implantation of the adjusting ring. The sled shaped nose portion on the end 52 of the dissecting ring causes the end 52 to be continuously biased upwardly sufficiently to maintain the ring 52 in a path in the anterior lamellae of the stroma. The magnetic force between the holder 72 and the dissecting ring also causes the dissecting ring to maintain its path within the anterior lamellae.

Rotation of the holding member 72 is continued with the pin 78 and magnetic attraction between the ring 51 and holder 72 serving to drive the ring about the cornea until the end portion 52 reaches the incision 83. At this time, a second incision (Fig. 9) is made perpendicular to the first incision, with the second incision being approximately 1 mm long. In any event the second incision extends from a position just above the hole 57 in the dissecting ring to a normal intersection with the first incision 83. The leading end 48 of the adjusting ring 47 is then brought into position with its tip end adjacent the tip 52 of the dissecting ring now lying below the corneal surface. As shown in Fig. 7, the clip member or link 61 is now placed with its leg portions 62 projecting into the holes 56 and 57 in the respective adjusting and dissecting rings and the body portion 63 of clip 61 is passed through the second incision 82 into position in the grooves 58 in the top side of the respective rings.



Next the holding member 72 is rotated in an opposite direction pulling the leading end 48 of the adjusting ring through the incision 83 and around the circular path previously made by the insertion of the dissecting ring.

5 The dissecting ring is moved in reverse rotation with the member 72 by means of the magnetic force between the holder 72 and ring 51. The plastic adjusting ring 47 is pulled by the metal dissecting ring. As the adjusting ring 47 is drawn into the cornea, the dissecting ring is being  
10 progressively withdrawn or "backed-out" until the leading end 48 of the adjusting ring is brought circularly back around to the incision 83, at which time the dissecting ring is fully withdrawn. The link 61 is then removed from the holes 56, 57 and slots 58 to release the connection between the dissecting and adjusting rings. The tip  
15 ends of the split ends 48, 49 of the adjusting ring are then grasped, such as by the ends of an adjustable caliper, and adjusted longitudinally endwise relative to one another to adjust the diameter of the ring and thereby bring the  
20 shape of the cornea into coincidence with the indicia 81 on the target image (Fig. 11). Also shown in Fig. 11 is an image of somewhat concentric circles 83 similar to that projected from a viewing surface by a corneoscope. The rings are produced by reflecting light from placido  
25 rings onto the present corneal surface being worked on. The distances from placido rings to cornea to image surface is the same as for constructing the target image. The concentricity and spacings of the rings 83 represents the topography of the eye being worked on. By superimposing  
30 the rings 83 onto the target image while adjustment of the ends of the adjusting ring 47 is taking place, perfect correction of the corneal shape may be affected by manipulating the ends of the adjusting ring until the indicia 81 and circles 83 are brought into coincidence.

The ends 48, 49 of the adjusting ring 47 are then fixed together such as by gluing or the like to permanently fix the correct shape of the cornea.

5           While particular embodiments of the present invention have been shown and described, it is apparent that changes and modifications may be made without departing from this invention in its broader aspects, and therefore, the aim  
10           in the appended claims is to cover all such changes and modifications as fall within the true spirit and scope  
          of the invention.

CLAIMS

1. An apparatus suitable for implantation into the cornea of the eye to permit adjustment of the corneal shape, characterised by: a ring shaped adjusting member (47) having separable end portions (48, 49) and means on one of the separable end portions capable of facilitating movement of said member (47) through corneal tissue.
2. An apparatus as claimed in claim 1 characterised in that the said ring shaped member (47) is made from a flexible material which is capable of acceptance by corneal tissue.
3. An apparatus as claimed in claim 1 or 2, characterised in that said separable end portions are arranged so that the one end is insertable into the cornea and is movable in a horizontal circular path therethrough whilst the trailing other end portion is still outside the anterior surface of the cornea, said movement facilitating means including an asymmetrically rounded portion which is sled-shaped to apply a biasing force to the member transverse to its normal path of movement within the cornea.
4. An apparatus as claimed in any one of claims 1 to 3 characterised in that the sectional shape of the adjusting member (47) is ellipsoid.
5. An apparatus as claimed in any one of claims 1 to 4 characterised in that one of the separable end portions has means thereon for connecting said adjusting member to an insertion device.
6. An apparatus as claimed in claim 5 characterised in that said connecting means includes a hole (57) near the end of said one separable end portion and a longitudinal slot (58) in the outer surface of said adjusting member (47) connecting

said hole to the tip of said one separable end portion.

5 7. An apparatus for implantation within the stroma of the cornea of the eye to permit adjustment of the corneal shape characterised by; a split ring shaped adjusting member (47); said ring shaped adjusting member being ovaloid in cross sectional shape and arranged so that its major cross sectional axis is alignable with the slope of a corneal arc passing through the anterior pole of the cornea when implanted  
10 in a cornea.

8. An apparatus as claimed in claim 7 characterised in that the split ring shaped adjusting member (47) has a first end portion which is shaped to facilitate its  
15 insertion and movement within the corneal tissue.

9. An apparatus as claimed in claim 7 or 8 characterised in that said first end portion (48) is asymmetrically rounded such that, when implanted in the cornea the first end portion (48) is capable of being biased laterally  
20 toward the anterior of the stroma as it is moved within the cornea.

10. An apparatus suitable for inserting an adjusting ring into the anterior of the cornea for making adjustments in  
25 the shape of the cornea characterised by a ring shaped cutting member (51) capable of being inserted into the cornea of the eye and having overlapping end portions (52, 53); one end (52) of said ring shaped cutting member (51) being substantially rounded to facilitate, in use, its easy  
30 movement along a circular path in the tissue of the cornea, means (56) near said one end (52) of said ring shaped cutting member (51) for releasably attaching one end of an adjusting ring (47).

11. An apparatus as claimed in claim 10 wherein said one end (52) of said ring shaped cutting member (51) is asymmetrically rounded in a sled shape to bias said ring shaped cutting member (51) in a direction which is transverse, in use, to its path of movement as it moves within the corneal tissue.

12. An apparatus as claimed in claim 10 or 11 characterised in that said ring shaped member (51) is substantially circular in cross sectional shape.

13. An apparatus as claimed in any one of claims 10 to 12 characterised in that holding means (71) are provided for holding said cutting member (51).

14. An apparatus as claimed in claim 13 wherein said cutting member (51) has magnetic properties and said holding means (71) includes a magnet means (72) for applying a holding force to said cutting member (51).

15. An apparatus as claimed in claim 14 characterised in that said holding means (71) has a concave surface (76) sized such as, in use, to receive the convex outer surface of the cornea of the eye.

16. An apparatus as claimed in claim 15 characterised in that a circular groove (77) is concentrically formed within said concave surface (76) for receiving said ring shaped cutting member (51) and for maintaining, in use, said ring shaped cutting member (51) in a circular path as it is being moved through the corneal tissue.

17. An apparatus as claimed in claim 16 characterised in that means (78) are provided in said groove (77) for engaging the other end of the cutting member (51) to facilitate, in use, movement of said cutting member (51)

into and through the corneal tissue.

18. An apparatus as claimed in claim 17 wherein the other end (54) of said cutting member (51) is blunt  
5 to facilitate its engagement with the engaging means (78) on said holding means (71).

19. An apparatus as claimed in any one of claims 10 to 18 characterised in that said releasably attaching means  
10 includes a hole (56) near said one end of said cutting member (51) and a longitudinal slot (58) in the upper-facing surface of said cutting member (51) extending from said hole (57) to the tip end of said one end of  
said cutting member (52)

20. An apparatus for inserting an adjusting ring (47) into the corneal tissue of the eye characterised by; an  
15 elongated member (72); a concave concentric surface (76) on one end of said elongated member (72); surface means (73) on at least a portion of said elongated member (72) to  
20 facilitate rotation of said elongated member (72); and magnet means for generating a magnetic field near said one end of said elongated member (72).

21. An apparatus as claimed in claim 20 characterised  
25 in that concentric groove means (77) are formed in said concave surface (76).

22. An apparatus as claimed in claim 21 wherein said groove  
30 means (77) is arranged to receive a split dissecting ring (51) and further including pin means (78) extending outwardly from said groove (77) for engaging one end of said dissecting ring (51).

23. An apparatus as claimed in claim 22 characterised in that said elongated member (72) is cylindrical and further including circular sleeve means (74) arranged for receiving said elongated member (72) therein.

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24. A system for implanting an adjusting ring (47) into the corneal tissue of the eye for permitting adjusting of the shape of the cornea; characterised by ring shaped dissecting means (51) having split end portions (52, 53) to permit, in use, insertion of one end portion (52) into the tissue of the cornea while the other trailing end portion (53) is outside the anterior surface of the cornea; means for holding said dissecting means (72) for movement within the cornea; ring shaped adjusting means (47) having split end portions (48, 49) to permit insertion of one end portion into the corneal tissue while the other trailing end portion is outside the anterior surface of the cornea; and means for releasably connecting (56, 57, 61, 62) said respective one end portions (48, 49).

20

25. A system as claimed in claim 24 characterised in that said ring-shaped adjusting means (47) is ellipsoid in cross-section.

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26. A system as claimed in claim 25 characterised in that the major axis of said ellipsoid shaped adjusting means (47) is alignable, when inserted within a cornea, with the slope of a corneal arc passing through the anterior pole of the cornea.

30

27. A system as claimed in any one of claims 24 to 26 characterised in that said respective one end portions (48, 49) each have an asymmetrically rounded tip end for generating a transverse biasing force to the end portion when, in use, it is moved in an inserting direction within

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corneal tissue.

28. A system as claimed in any one of claims 24 to 27 characterised in that each of said respective one end portions (48, 49) have holes (56, 57) near their tip ends (48, 49) for connecting said respective one end portions.

29. A system as claimed in claim 28 characterised in that removable clip means (61, 62) are provided for insertion into said holes (56, 57) for holding said dissecting means (51) and adjusting means (47) from relative longitudinal movement.

30. A system as claimed in claim 29 characterised in that longitudinal slot means (58) are formed in the upper surface of said adjusting means (47) and dissecting means (51), said slot means extending from said holes to the tip end of the respective one end portions (48, 49).





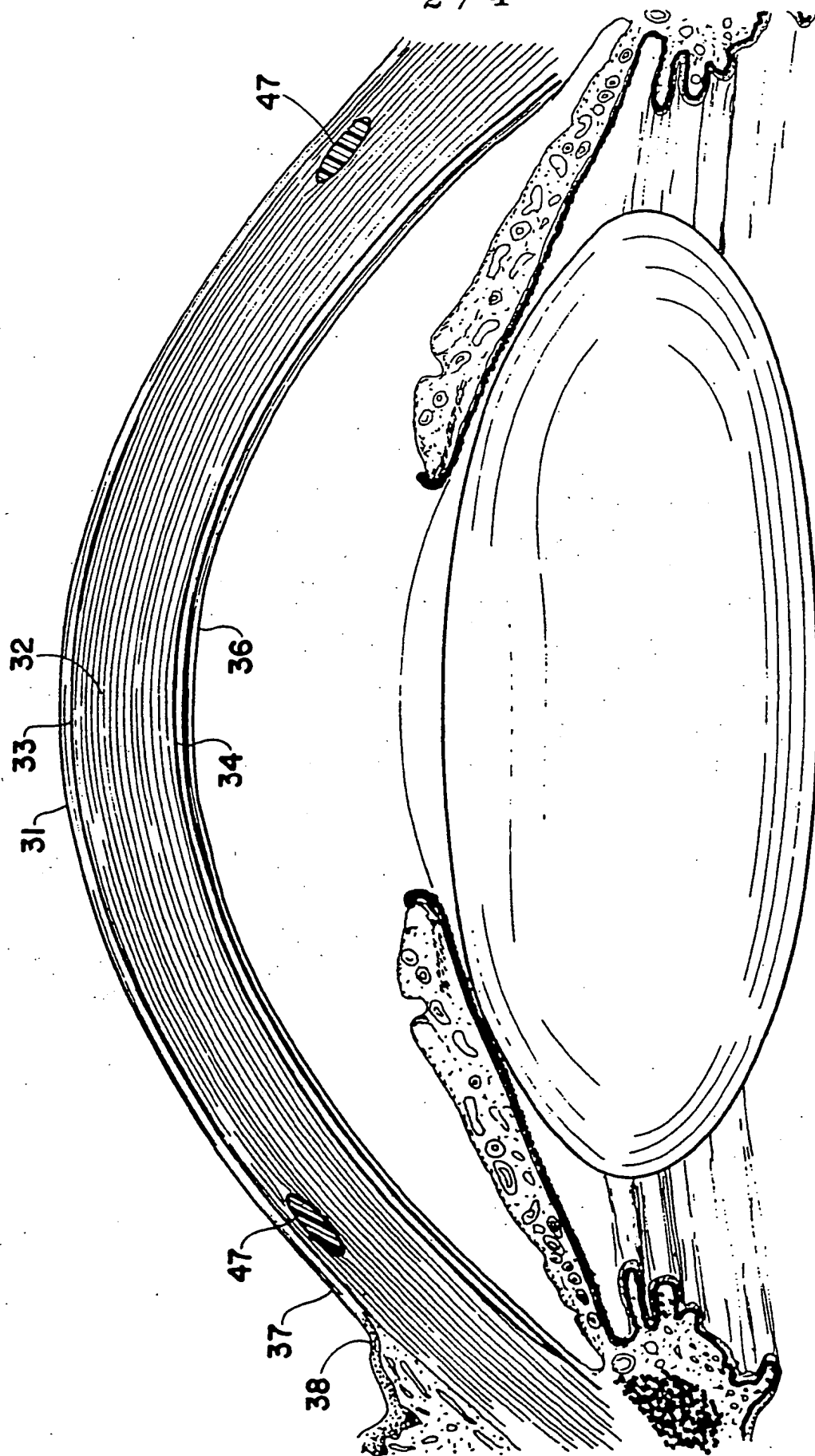
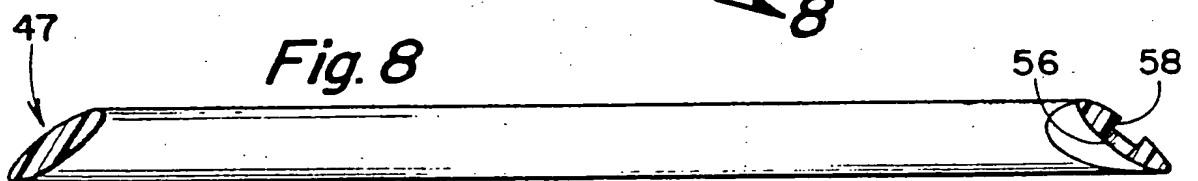
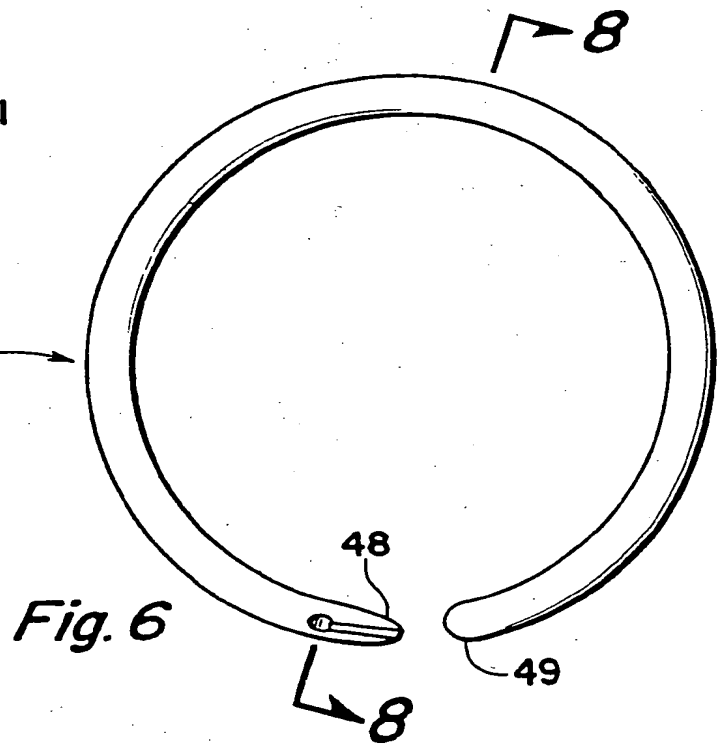
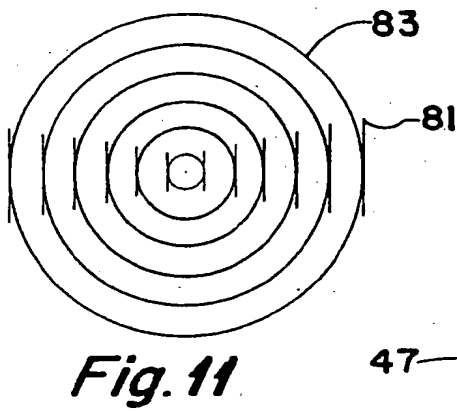
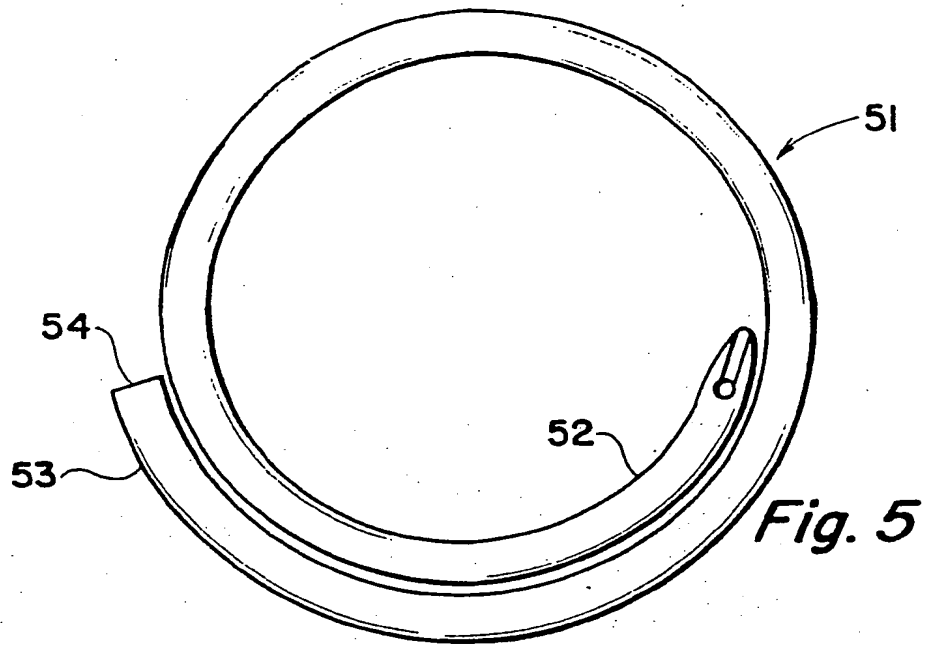
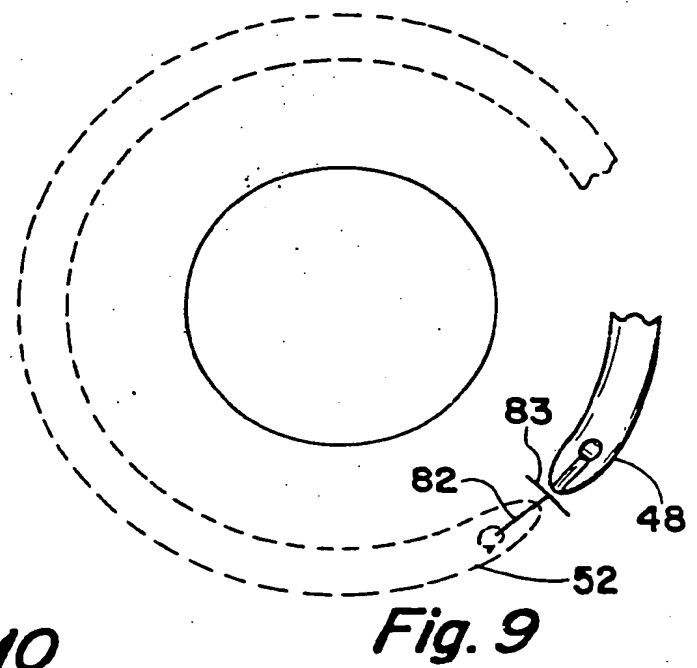
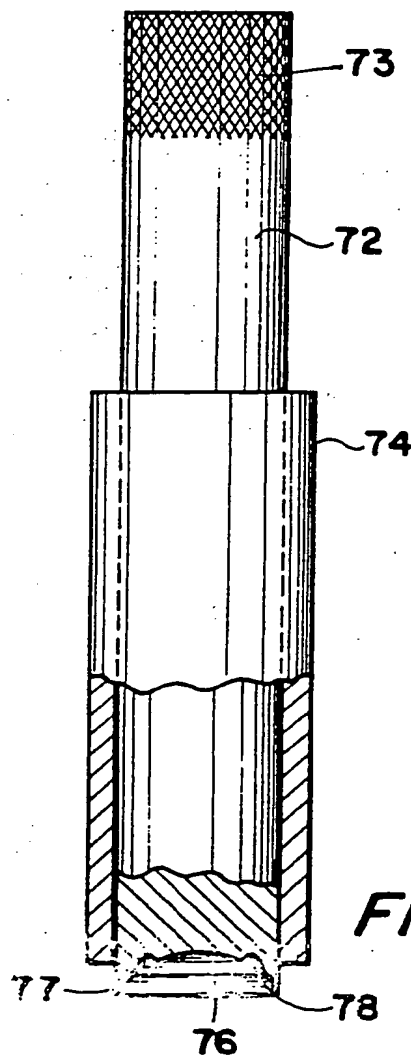
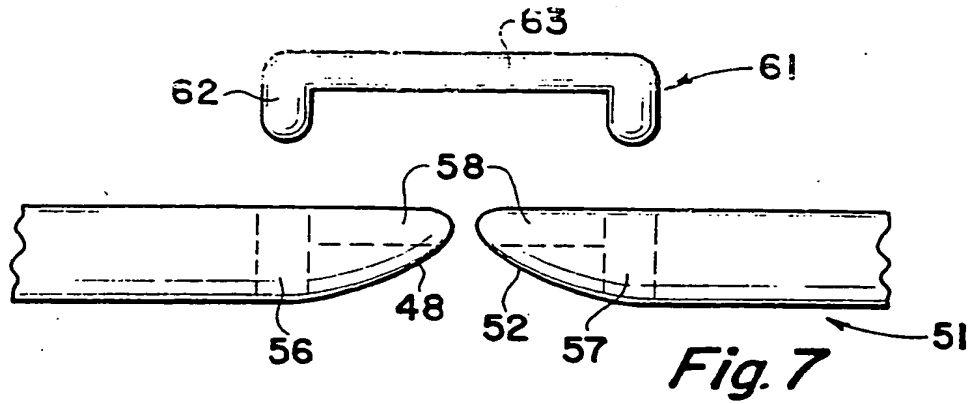


Fig. 4







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# EUROPEAN SEARCH REPORT

0083494

Application number

EP 82 30 6834

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 7)
A	US-A-3 064 643 (J.H.DIXON) *The whole document*	1,2	A 61 F 9/00
A	FR-A-2 354 752 (C.M.L.LECOCQ) *Claim 1*	1,2	
A	US-A-2 754 520 (J.H.CRAWFORD) *The whole document*	1-3,7-11	
A	US-A-4 053 953 (L.FLOM & K.J.RODGERSON) *Column 17, lines 7-33; column 22, lines 19-52; figures 34-37,40*	1-7,13,22,23	
A,P	GB-A-2 095 119 (J.L.TENNANT & H.J.SMIRMAUL) *The whole document*	1,2	TECHNICAL FIELDS SEARCHED (Int. Cl. 7) A 61 F A 61 B
A,P	US-A-4 346 482 (J.L.TENNANT AND H.J.SMIRMAUL) *The whole document*	1	
A	DE-B-2 719 688 (W.HENNING) *Claim 1*	13,14	
A	US-A-3 656 481 (R.A.NESS) *Abstract*	13,14	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 05-04-1983	Examiner WOLF C.H.S.
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons &amp; : member of the same patent family, corresponding document</p>			

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A	US-A-2 818 866 (W.C.THOMAS) *Figure 4; column 3, lines 1-9*	3,10	
A	GB-A-2 029 235 (R.P.JENSEN) *Figure 1*	28,29	
			TECHNICAL FIELDS SEARCHED (Int. Cl. 3)
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Place of search THE HAGUE		Date of completion of the search 05-04-1983	Examiner WOLF C.H.S.
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